## In the Claims:

Please amend the claims as follows:

## 1-10. (Cancelled)

- 11. (New) A population of human marrow stromal cells enriched for small and rapidly self-renewing stem (RS) cells, said population comprising about 95% RS cells, wherein said RS cells are about seven microns in diameter, comprise granular and agranular cells, and do not express STRO-1, PDGF-R, EGF-R, CD10 and CD147.
- 12. (New) The population of human marrow stromal cells of claim 11, wherein said population enriched for RS cells has an increased capacity for multilineage differentiation compared to a population of human marrow stromal cells not enriched for RS cells.
- 13. (New) The population of human marrow stromal cells of claim 11, wherein said RS cells have a high nucleus to cytoplasm ratio.
- 14. (New) The population of human marrow stromal cells of claim 11, further wherein said RS cells express at least one of the polypeptides selected from the group consisting of VEGF receptor-2 (FLK-1), TRK (an NGF receptor), transferrin receptor, annexin II (lipocortin 2) and CD49e (integrin alpha 5).
- 15. (New) The population of human marrow stromal cells of claim 11, further wherein cells within a first subtype of said RS cells are agranular and express VEGF receptor-2 (FLK-1), TRK (an NGF receptor), transferrin receptor, and annexin II.
- 16. (New) The population of human marrow stromal cells of claim 15, further wherein said cells within a first subtype express CD44 (Hyaluronic acid receptor), CD49e (integrin alpha 5) and CD59.

2

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- 17. (New) The population of human marrow stromal cells of claim 11, further wherein cells within a second subtype of said RS cells are granular, express CD49e (integrin alpha 5) and do not express CD44 (Hyaluronic acid receptor) and CD59.
- 18. (New) The population of human marrow stromal cells of claim 17, further wherein said cells within a second subtype do not express CD81 and CD90.
- 19. (New) The population of human marrow stromal cells of claim 11, further wherein said RS cells express heat shock protein-27, tumor rejection antigen, glutathione-S transferase, peroxiredoxin 1, voltage-dependent-anion channel-2, protein kinase C substrate, phosphatase 2A inhibitor, esterase D, RNase A, initiation factor 5a, elongation factor 1-alpha, ribosomal protein S12, ribosomal protein large P1, ribosomal protein large P2, transcription factor BTF 3a, annexin I, destrin, myosin light chain, lactate dehydrogenase A, glycerolaldehyde-3-P dehydrogenase, citrate synthetase, transketolase, P-glycerolmutase, aldoketo reductase 7(A2), alpha-amylase inhibitor CM3, enoyl-CoA hydratase, and proteosome subunit alpha-4.